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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/699,051	10/30/2003	Brian R. Reynolds	1001.1716101	1188
28075 7590 08/20/2008 CROMPTON, SEAGER & TUFTE, LLC 1221 NICOLLET AVENUE			EXAMINER	
			HOEKSTRA, JEFFREY GERBEN	
SUITE 800 MINNEAPOLIS, MN 55403-2420			ART UNIT	PAPER NUMBER
			3736	
			MAIL DATE	DELIVERY MODE
			08/20/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)	
	10/699,051	REYNOLDS ET AL.	
Office Action Summary	Examiner	Art Unit	
	JEFFREY G. HOEKSTRA	3736	
The MAILING DATE of this communication ap Period for Reply	ppears on the cover sheet with the c	correspondence address	
A SHORTENED STATUTORY PERIOD FOR REPI WHICHEVER IS LONGER, FROM THE MAILING I - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period - Failure to reply within the set or extended period for reply will, by statu Any reply received by the Office later than three months after the maili earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION .136(a). In no event, however, may a reply be tird will apply and will expire SIX (6) MONTHS from te, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).	
Status			
Responsive to communication(s) filed on <u>07</u> 2a) This action is FINAL . 2b) Th Since this application is in condition for allowed closed in accordance with the practice under	is action is non-final. ance except for formal matters, pro		
Disposition of Claims			
4) Claim(s) 1-12,21,22 and 24-34 is/are pending 4a) Of the above claim(s) is/are withdra 5) Claim(s) is/are allowed. 6) Claim(s) 1-12,21,22 and 24-34 is/are rejected 7) Claim(s) is/are objected to. 8) Claim(s) are subject to restriction and/	awn from consideration.		
Application Papers			
9) ☐ The specification is objected to by the Examin 10) ☑ The drawing(s) filed on 30 October 2003 is/ar Applicant may not request that any objection to the Replacement drawing sheet(s) including the corre 11) ☐ The oath or declaration is objected to by the E	e: a)⊠ accepted or b)⊡ objected e drawing(s) be held in abeyance. Sec ction is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreig a) All b) Some * c) None of: 1. Certified copies of the priority documer 2. Certified copies of the priority documer 3. Copies of the certified copies of the pri application from the International Bures * See the attached detailed Office action for a list	nts have been received. nts have been received in Applicati ority documents have been receive au (PCT Rule 17.2(a)).	ion No ed in this National Stage	
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail D: 5) Notice of Informal F 6) Other:	ate	

DETAILED ACTION

Continued Examination Under 37 CFR 1.114

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 08/07/2008 has been entered.

Notice of Amendment

2. In response to the amendment filed on 08/07/2008, amended claim(s) 1, 3, 4, 7-9, 21, and 22 and new claim(s) 32-34 is/are acknowledged. The current rejections of the claim(s) 1-12, 21, 22, and 24-31 is/are *withdrawn*. The following new and reiterated grounds of rejection are set forth:

Claim Objections

3. Claims 1 and 3 are objected to because of the following informalities: the first positive recitations of "the final medical device" appear to lack antecedent basis and may render the claims indefinite. Appropriate correction is required. The Examiner notes Applicant may have intended to positively recite "a final medical device", "the medical device", or the like.

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Claim Rejections - 35 USC § 102

4. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

- 5. Claims 1-12, 21, 22, and 24-34 are rejected under 35 U.S.C. 102(e) as being anticipated by Sharrow (US 2004/0167438 A1).
- 6. The Examiner notes Sharrow (US 2004/0167438 A1) incorporates by reference (Sharrow paragraphs 24, 34, and 41), the following:
- Zhou et al. (US 2004/0143239 A1, hereinafter Zhou),
- Reynolds et al. (US 2004/0167441 A1, hereinafter Reynolds 1),
- Nguyen et al. (US 5,772,609, hereinafter Nguyen),
- Palermo (US 6,139,510),
- Skujins et al. (US 6,918,882 B2, hereinafter Skujins), and
- Reynolds et al. (US 7,074,197 B2, hereinafter Reynolds 2).
- 7. For claims 1-12, 21, 22, and 24-34, Sharrow discloses a method for manufacturing an intracorporeal medical device (10), comprising the steps of:
 - providing an elongate core member (14) having a proximal region (16) and a distal region (18) (paragraphs 16-28) (as best seen in Figure 2);

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distally and/or proximally disposing a smooth thermoplastic polymer jacket (20) over said elongate core member (paragraphs 29-31) (as best seen in Figure 2), the polymer jacket having a substantially smooth outer surface (paragraphs 29-31);

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- winding in tension a coil (12) (the conventional winding technique as positively recited in paragraph 36) including fluorocarbon materials (the high performance PTFE polymer positively recited in paragraphs 30 and 36) (see specification page 7 lines 1-2) over the outer surface of said polymer jacket (paragraphs 32-39), wherein the coil includes an outer fluorocarbon-containing coating (paragraphs 32-39) (the high performance PTFE polymer positively recited in paragraphs 30 and 36) (see specification page 7 lines 1-2), wherein the coil includes a central metallic core material and an outer coating surrounding the central core material (paragraphs 35, 36, and 39) (Zhou paragraphs 29-31, 38, and 56);
- heating said polymer jacket (paragraphs 36-38), relieving the tension within the coil is relieve and the coil (paragraphs 36-38), and embedding the coil in said polymer jacket (paragraphs 36-38), wherein in the heating/tension-relieving/embedding the coil moves radially inward into the polymer jacket and a portion of the outer surface of the polymer jacket wicks outward between the adjacent windings of the coil (paragraphs 36-38), thereby altering the shape of the outer surface of the polymer jacket (paragraphs 36-38) and providing an outer surface of the polymer jacket relative to the coil that has desirable flexibility

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characteristics in the intracorporeal medical device (paragraphs 2, 3, and 37); and

- disposing a covering (22) over said jacket/coil member (paragraph 41),
- wherein the intracorporeal medical device is manufactured to include an
 outermost surface (412a) having a helical ridge extending around a
 circumference of the outermost surface formed at least in part by the coil (as best
 seen in Figure 6) (paragraphs 46-47).

Response to Arguments

- 8. Applicant's arguments with respect to claims 1-12, 21, 22, and 24-34 have been considered but are moot in view of the new ground(s) of rejection, wherein the new ground(s) of rejection relies upon a different interpretation of previously applied prior art applied to amended and new claims. However in the interest of advancing prosecution and in lieu of previously applied prior art, the Examiner notes the following:
- 9. Applicant's arguments filed 08/07/2008 have been fully considered but they are not persuasive. Applicant argues the anticipatory rejection of the claims under Sharrow, specifically arguing Sharrow does not disclose, teach, and/or fairly suggest the following: (a) the intracorporeal medical device is manufactured to include an outermost surface having a helical ridge extending around a circumference of the outermost surface formed at least in part by the coil or (b) the central core material is a metallic material. The Examiner disagrees, maintains the rejection as set forth above, and notes in response the following:

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10. In response to Applicant's argument (a), the Examiner notes Sharrow discloses the intracorporeal medical device is manufactured to include an outermost surface (412a) having a helical ridge extending around a circumference of the outermost surface formed at least in part by the coil (as best seen in Figure 6) (paragraphs 46-47).

11. In response to Applicant's argument (b), the Examiner notes Sharrow discloses the coil includes a central metallic core material and an outer coating surrounding the central metallic core material (paragraphs 35, 36, and 39) (Zhou paragraphs 29-31, 38, and 56).

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to JEFFREY G. HOEKSTRA whose telephone number is (571)272-7232. The examiner can normally be reached on Monday through Friday 8am to 5pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Max Hindenburg can be reached on (571)272-4726. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

/J.H./
Jeff Hoekstra
Examiner, Art Unit 3736

/Max Hindenburg/ Supervisory Patent Examiner, Art Unit 3736